

 Date of Report
 1/7/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. KRISHAMA			Patient ID		012206300073
Birthday		14 07 1993	Sample ID		11570335
Age at term		29.05	Sample Date		29/6/2022
Gestational age		12+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	45.8 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+2
PAPP-A	$2.45~\mathrm{mIU/ml}$	0.43	Method		CRL (<>Robinson)
fb-hCG	113.5 ng/ml	2.59	Scan date		28/06/2022
Risks at sampling date			Nuchal translucency 4		
Age Risk	ge Risk 1:683		Nuchal translucency MoM 2.69		
Biochemical T21 risk		1:53	Nasal bone Preser		Present
Combined trisomy 21 risk		>1:50			
Trisomy 13/18 + NT		>1:50			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is >1:50 , which indicates a increased risk			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a above the risk. After the result of the Trisomy 21 with NT test it is expected that among less than 50 women with the same data. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		